

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
9 January 2003 (09.01.2003)

PCT

(10) International Publication Number
WO 03/002131 A1

(51) International Patent Classification⁷: **A61K 35/74**,
C12N 1/20, A61P 31/04, 1/00

SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(21) International Application Number: PCT/EE02/00006

(22) International Filing Date: 21 June 2002 (21.06.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
P200100356 29 June 2001 (29.06.2001) EE

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(81) Designated States (national): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, GR, GU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MY, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,

(84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent
(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG).

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FI, GB, GD, GE, GH, GM, GR, GU, ID, IL, IN, IS, JP, KE,
KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG,
MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU,
SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG,
UZ, VN, YU, ZA, ZM, ZW, ARIPO patent (GH, GM, KE, LS,
MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent
(AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent
(AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU,
MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI,
CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)
- of inventorship (Rule 4.17(iv)) for US only

Published:

- with international search report
- with (an) indication(s) in relation to deposited biological
material furnished under Rule 13bis separately from the
description

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: STRAIN OF MICRO-ORGANISM LACTOBACILLUS FERMENTUM ME-3 AS NOVEL ANTI-MICROBIAL AND
ANTIOXIDATIVE PROBIOTIC

(57) Abstract: The strain of micro-organism *Lactobacillus fermentum* ME-3 is a novel anti-microbial and anti-oxidative probiotic. It has a high anti-microbial effect on *Escherichia coli*, *Shigella sonnei*, *Staphylococcus aureus*, *Salmonella typhimurium*, and moderate activity against *Helicobacter pylori* strains. The strain of micro-organism possesses Mn-superoxide dismutase and both its lysates and intact cells have high anti-oxidative activity, increasing the glutathione red-ox ratio in blood sera and able to capture toxic hydroxyl radicals. The strain of micro-organism could be used as a probiotic for the production of functional food (yoghurt, cheese) and non-comestibles (tablets, capsules) for the prophylaxis of intestinal and uroinfections, both for the prevention and treatment of chronic diseases, caused by prolonged oxidative stress.

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STRAIN OF MICRO-ORGANISM *LACTOBACILLUS FERMENTUM* ME-3 AS
NOVEL ANTI-MICROBIAL AND ANTI-OXIDATIVE PROBIOTIC

TECHNICAL FIELD

5 The present invention relates to biotechnology and will be used as a novel probiotic applied in the production of functional food (yoghurt, cheese) and non-food preparations (tablets, capsules) for the prevention or treatment of different diseases.

10

BACKGROUND ART

Probiotics are living microbial food additives, that have beneficial effect on the microbiological balance of the intestine and human health. Probiotics are used as functional
15 food. Functional food is foodstuff, consumed additionally to usual food and containing bio-preparations (incl. probiotics) or other components favourably influencing human health or decreasing disease risks.

Probiotics are consumed as components of food (probiotic
20 yoghurt or cheese) or non-food preparations (lyophilised microbial cultures).

Most of probiotics are lactic acid bacteria, mainly lactobacilli. Lactobacilli are non-pathogenic micro-
25 organisms, colonising the human intestinal and urogenital tract from early childhood to old age. Nowadays, several commercial probiotic lactobacilli are successfully used, among which *Lactobacillus rhamnosus* GG (Saxelin M. *Lactobacillus* GG - a human probiotic strain with thorough
30 clinical documentation. Food Rev Int 1997; 13:293-313) is the best known. Recently some new strains of lactobacilli have been described and patented, for example *L. reuterii* (Korea patent KR211529, C12N 1/20, Korea Institute Science Technology, 1999), isolated from animal organism and for this
35 reason inappropriate for human usage.

Several strains of *Lactobacillus fermentum* are used for correction and stabilisation of intestinal micro-flora in case of dysbacterioses and urogenital infections with different ethiologies. The strain of micro-organism
5 *Lactobacillus fermentum* 39 is used for producing the bacterial biological preparation (PCT/SU89/00264 (WO 91/05852), C12N 1/20, A61K 35/74, University of Tartu, 1991). The strain *Lactobacillus fermentum* 90-TS-4 (RU2133272, C12N 1/20, A61K 35/74, Akivo Lentsner et al., 1999) is
10 characterised by lectin typing as a mannose-sensitive profile of the cell wall. The preparation is prescribed for use in gynaecology.

There are some well-known probiotics, targeted against only
15 one pathogen (for example Salmonella) (US5478557, A61K 35/74, US Agriculture, 1995; US5340577, A61K 35/74, US Army, 1994). Up to the present no strain of lactobacilli with an extensive anti-microbial effect against numerous pathogens and opportunistic pathogens has been described.

20 Likewise, yet no such strain of micro-organism is known that could have natural antibiotic resistance against drugs most frequently used in the treatment of infections. This property would permit to use such strains in case of antibiotic-
25 treated patients. A set of different micro-organisms is used in veterinary, containing also one strain of *L. fermentum*, but this strain does not have a concurrent anti-microbial and anti-oxidative effect (RU2119796, A61K 35/66, Zakrõtoje aktsionernoje obshestvo "BAKS", 1998).

30 Anti-oxidative preparations like vitamin E and C, betacarotene a.o. nowadays get much attention in connection to healthy nutrition. Excessive formation of reactive oxygen species (ROS) in tissue respiration can cause the damage of cells and the course of tissues. The formation of active

oxygen may depend on some stress factors, such as alcohol, peroxides, and some drugs.

Usually excessive oxidation is closely connected with nutritional diseases, age, arteriosclerosis, misfunctions of the central nervous system and the intestinal tract, cancer a.o. pathological conditions. An organism has several defence systems against the toxigenicity of oxygen. It is important to take anti-oxidative substances to guarantee the functioning of these systems.

Of known solutions, the closest to this invention is the patent describing anti-oxidative food, an anti-oxidative preparation and a method of antioxidation (EPO649603, A23L 3/3472, A23L 3/3571, Otsuka Pharma Co Ltd. 1995). The object of this invention is a preparation that contains a natural substance involving mangan (leaves of tea plant) and the micro-organism *Lactobacillus plantarum* that produces catalase and a superoxidase-dismutase system, thus increasing the anti-oxidative activity of the host organism. The authors of this invention declare that the preparation prevents diseases developing due to active oxygen. However, they do not describe the effect of particular *Lactobacillus* strain with a decreasing anti-oxidative activity or capturing hydroxyl radicals *in vitro*. Besides that, this strain of micro-organism is also imperfect because for getting the presumable anti-oxidative effect in an organism (*in vivo*), it is necessary to add some Mn-containing raw material (leaves of tea plant) to the preparation, because it is only in this case its SOD (superoxide dismutase) activity is realised.

DISCLOSURE OF THE INVENTION

The aim of this invention is to present the strain of micro-organism as a novel anti-microbial and anti-oxidative probiotic for use in pharmaceutical and food industry, also

in medicine as an antibiotic-resistant preparation for the prophylaxis and treatment of gastrointestinal and uroinfections, and against oxidative stress.

- 5 The object of the investigation - the strain of microorganism *Lactobacillus fermentum* ME-3 was isolated from a faecal sample of a healthy child during a comparative study of the micro-flora of Estonian and Swedish children, using MRS (Oxoid) media and cultivating it in a CO₂ environment (Sepp
10 et al., Intestinal microflora of Estonian and Swedish infants, Acta Paediatrica, 1997, 86, 956-961).

The strain of micro-organism *Lactobacillus fermentum* ME-3 was isolated by seeding the dilutions of the faeces of healthy
15 one-year-old Estonian child (10^{-2} - 10^7 in phosphate buffer with 0.04% thioglycol acid; pH 7.2). The dilutions were seeded on freshly prepared MRS agar-media and cultivated at 37°C in a CO₂ environment. The strain, which is the object of
20 invention, was isolated from a 10^{-5} dilution on the basis of the characteristic morphology of colonies and cells. A provisional and more precise identification followed as described next. Using additional tests, the strain was
selected from other lactobacilli isolated from the same child on the basis of its special properties.

25

The fact that the microbial strain *Lactobacillus fermentum* ME-3 originates from the intestinal tract of a healthy child
proves its GRAS (*generally recognised as safe*) status, i.e. that this strain of micro-organism is harmless for human
30 organism and it is suitable for oral application.

Cultural-morphological characteristics were determined after cultivating the strain on MRS agar and in MRS broth media (OXOID). Microbial cells are Gram-positive rods of regular

shape located in parallel chains, nonspore, of medium thickness and different length (2x 3-5 μ m).

Physiological-biochemical characteristics: MRS broth was suitable for cultivating the microbial strain during 24-48 hours in a 10% CO₂ environment, after which homogeneous turbid growth occurred in the broth. The colonies of micro-organism on MRS agar are white, rounded, with a regular edging.

The optimal growth temperature is 37°C, it multiplies also at 45°C, but it does not grow at 15°C. The optimal growth environment is at pH 6.5.

The negative catalase test, gas production by fermentation of glucose, production of NH₃ from arginine, and lysozyme production are the main properties. During reproduction in milk it produces 1,07% of acid.

The strain with above-mentioned characteristics was identified on the basis of biochemical activity with API 50 CHL System (BioMerieux, France) kit as *Lactobacillus fermentum* (ID% 99.6, T 0.87, only 1 test contra). The following sugars and alcohols were fermented - ribose, galactose, D-glucose, D-fructose, D-mannose, esculine, maltose, lactose, melibiose, saccharose, D-raffinose, D-tagatose and gluconate.

The profile of the metabolites of *Lactobacillus fermentum* ME-3 was characteristic of heterofermentative metabolism, determined by the gas chromatographic method (Hewlett-Packard model 6890). The profile of fermentation depended on environment of incubation: besides lactic and acetic acids a big amount of succinic acid was produced in a CO₂ environment, but in an anaerobic environment much of ethanol was produced in addition to the above-mentioned substances (Table 1). Both succinic acid and ethanol can strengthen the

stable properties of the microbial strain in milk fermented by this strain.

Table 1. The concentration of acetic acid, lactic acid, succinic acid and ethanol (mg/ml) in MRS media in cultivation of *Lactobacillus fermentum* ME-3 in microaerophilic and anaerobic environment during 24 and 48 h.

<i>Lactobacillus fermentum</i> ME-3	Lactic acid		Acetic acid		Succinic acid		Ethanol	
	24 h	48 h	24 h	48 h	24 h	48 h	24 h	48 h
CO ₂ environment	10.6	11.1	0.8	0.9	18.4	19.5	9.8	7.5
Anaerobic environment	8.2	8.8	1.0	1.0	5.7	9.7	7	33.3

10 Molecular identification.

Molecular identification by ITS-PCR (internal transcribed spacer - polymerase chain reaction) using *Lactobacillus fermentum* ATCC 14931 as the reference strain verified the previous identification with API 50 CHL.

15

The micro-organism with the above-mentioned properties was deposited in Deutsche Sammlung für Mikroorganismen und Zellkulturen GmbH-s, the registration number of the deposit is DSM 14241 (19.04.2001).

20

Anti-microbial activity

Lactobacillus fermentum ME-3 expresses a high anti-microbial effect on *Escherichia coli*, *Shigella sonnei*, *Staphylococcus aureus*, *Salmonella typhimurium* 1 and 2, and *Helicobacter pylori* strains in vitro (Table 2).

25

Table 2. Anti-microbial activity of strain *Lactobacillus fermentum* ME-3 on modified MRS-agar, in MRS broth and milk.

Lactobacillus fermentum ME-3	Escherichia coli	Shigella sonnei	Staphylococcus aureus	Salmonella typhimurium 1 and 2	Helicobacter pylori	
MRS-agar	Inhibition zone (mm)					
CO ₂ /anaerobic environment	24/22	26/21	20/19	25.8 / 24.7	23.8 / 19.7	13.2/13.1
MRS broth	Decrease of total count (log ₁₀) compared with initial count					
	log 6.0	Log 6.7	log 0.8	log 6.3	log 3.8	not determined
Milk	Suppression after different interval of time (24 - 48 h)					
	24 t	32 t	24 t	32 t	48 t	not determined

5

Using milk fermentation it was possible to show that pathogens inoculated into milk were killed in 24-48 h if milk was fermented with *Lactobacillus fermentum* ME-3. Such property of the strain could help to prevent the multiplication of pathogens in products (yoghurt, cheese) fermented by this strain, and prevent food infections. Organic acids and ethanol produced by *Lactobacillus fermentum* ME-3 could ensure the high anti-microbial effect of this microbe.

15

Resistance to antibiotics

According to a disk-diffusion test (BBL Sensi disks) and an E-test (AB Biodisk, Solna) *Lactobacillus fermentum* ME-3 was resistant to metronidazole, ofloxacin, aztreonam, cefoxitin

and TMP-SMX. This allows to use the strain *L. fermentum* ME-3 as a preparation accompanying antibiotic treatment in case of intestinal and uroinfections.

5 Surface structures of microbial cell

The carbohydrate profile of the surface structure of microbial cells of *Lactobacillus fermentum* ME-3 was determined by lectin typing. The strain of lactobacilli agglutinated with *Griffonia simplicifolia* I lectin, which is
10 specific to Gal and GalNAc ligands in the cell wall.

The strain *Lactobacillus fermentum* ME-3 did not react with the following other lectins: *Concanavalin ensiformis* (Con A), *Griffonia simplicifolia* II, *Arachis hypogaea* (PNA), *Vicia*
15 *sativa* (VSA) and *Tritium vulgare* (WGA).

Hence the special composition of the glycocalyx of the cell wall of *Lactobacillus* ME-3 became clear with lectin typing, it contained residues of galactose and N- acetyl-galactose-
20 amine. These compounds act as adhesins for engaging the receptors of mucosa on the epithelial cells of the upper urinary tract.

This is a possibility for blocking the mannose-resistant pili
25 of *Escherichia coli* that makes our strain applicable in the prophylaxis of urinary tract infections.

Anti-oxidative properties

Lactobacilli were incubated in a MRS broth (Oxoid Ltd.) for
30 24 h and centrifuged at 4° C (1500 p/min) 10 min for getting a precipitate, washed with isotonic salt (4°C) and suspended to the density of 1.15% KCl (Sigma, USA). The density of the suspension was at OD₆₀₀ 1.1 10⁹ bacterial cells in ml⁻¹.

To get lysates, the cells were disrupted by sonification (B-
35 12 Branson Sonic Power Company, Danbury, Connecticut) in 35

vibrations s^{-1} 10 min in an ice bath and then for 10 min at $-18^{\circ}C$. The suspension was centrifuged at $4^{\circ}C$ 10000 g/r for 10 min and the supernatant was filtered (MILLEY-GS, sterile, 0.22 μm ; Millipore S.A., 67 Molsheim, France) to get a cell-free extract. *Lactobacillus fermentum* ME-3 cells and lysate produced H_2O_2 in a remarkable amount (Table 3).

Table 3. Total anti-oxidative capacity of *Lactobacillus fermentum* strains ME-3 and E-338-1-1 (according to LA and TAS tests), hydrogen peroxide content, glutathione red-ox ratio and activity of superoxide dismutase.

Properties	<i>Lactobacillus fermentum</i> ME-3	<i>Lactobacillus fermentum</i> E-338-1-1
	Intact cells	Intact cells
TAA in LA-test (%)	29 ± 0.7 (n=5)	0
TAS (mmol/L)	0.16 ± 0.03 (n=5)	0
H_2O_2 ($\mu g/ml$)	31 ± 26 (n=3)	49 ± 20 (n=3)
	Lysate of cells	Lysate of cells
LA-test (%)	59 ± 3.8 (n=5)	0
H_2O_2 ($\mu g/ml$)	229 ± 37 (n=4)	137 ± 25 (n=3)
TGSH	12.5 ± 4.1	5.5 ± 3.0
GSSG ($\mu g/ml$)	2.59 ± 2.01	5.5 ± 2.4
GSH ($\mu g/ml$)	9.95 ± 3.30	Marks
GSSG/GSH	0.28 ± 0.17	0 ^e
SOD (U/mg protein)	0.859 ± 0.309 (n=3)	Not determined

Explanations: LA-test - linolenic acid test; TAA - total anti-oxidative activity; TAS - total anti-oxidative status; GSSG - oxidized glutathione; GSH - reduced glutathione;

GSSG/GSH - glutathione red-ox ratio; SOD - superoxide dismutase

Lactobacillus fermentum ME-3 has a Mn-SOD activity determined
5 by electrophoresis. For determining the SOD type *L. fermentum*
ME-3 cell-free extract (30 µg protein) was separated on 10%
not-denaturated polyamide-acrylic gel. SOD isoenzyme was
determined by influencing this gel with 15mM H₂O₂, after which
the SOD activity persisted. Explanation: H₂O₂ inhibits Fe-SOD,
10 but does not inhibit Mn-SOD. This proves that *Lactobacillus*
fermentum ME-3 has Mn-SOD activity.

The strain *Lactobacillus fermentum* ME-3 shows a high TAA
(total antioxidative activity) value in a lipid environment
15 on the basis of a linolenic acid test, also a high TAS (total
antioxidative status) value in a hydrate environment (Radox
kit, UK). In Table 3, data of the anti-oxidative strain
Lactobacillus fermentum E-338-1 is added for comparison
(Table 3).

20 The cells and lysates of the strain *Lactobacillus fermentum*
ME-3 catch hydroxyl radicals, this has been proven by the
terephthalic acid method (27% ± 5%). 15mM of reduced
glutathione was used for comparison as a well-known scavenger
25 of hydroxyl radicals (84±4,6%). *Lactobacillus fermentum* ME-3
survived in a highly oxidative H₂O₂ environment.

Re-cultivation of the lyophilised culture kept in room
temperature for a long time proved the viability of the
30 strain and the persistence of properties. This ensures that
the lyophilised strain of *Lactobacillus fermentum* ME-3 could
be used as a non-comestible product in a scheme of functional
food.

BEST MODE FOR CARRYING OUT THE INVENTION

An example of the preparation of a yoghurt with highly anti-oxidative properties based on the strain *Lactobacillus fermentum* ME-3 and the trial of consuming the yoghurt by healthy volunteers.

Lactobacillus fermentum ME-3 pure culture in 0.15% MRS-agar is used for producing the yoghurt, additionally the pure cultures of *Lactobacillus plantarum* and *Lactobacillus buchneri* are seeded into fresh goat milk autoclaved for 20 min at 110°C. Three cultures of these strains of lactobacilli are mixed in equal proportions together with 2% of *Streptococcus thermophilus* and are added in 0.2% of content into autoclaved goat milk.

Lactobacillus fermentum ME-3 with strains of lactobacilli and streptococci will guarantee tasty and highly anti-oxidative yoghurt (Table 4).

Table 4. The anti-oxidative activity of *Lactobacillus fermentum* ME-3 pure culture and probiotic yoghurt

Strain	Total anti-oxidative activity (TAA %)	
	Cells	Yoghurt
<i>Lactobacillus fermentum</i> ME-3	29	70

In tables 5 and 6, the changes of the intestinal micro-flora and indices of oxidative stress of blood sera of healthy volunteers are shown before and after taking the probiotic goat milk yoghurt during 3 weeks. These changes prove the anti-oxidative (incl. anti-atherogenic) effect on human organism.

Even a higher total anti-oxidative activity of goat milk yoghurt compared with the total anti-oxidative activity of intact microbial cells of *Lactobacillus fermentum* ME-3 is shown in table 5.

5

Additive microbial strains ensure the standard acidity and consistence of yoghurt.

Table 5. The changes of intestinal micro-flora of healthy volunteers (n=16) before and after consuming probiotic goat milk yoghurt during 3 weeks

	Before		After	
	Persons colonised	Lactoflora ratio (%)	Persons colonised	Lactoflora ratio (%)
Consuming goat milk yoghurt (n=16 persons)				
<i>L. fermentum</i>	4*	0,7 - 5,77	16*	0,5 - 49,9#
Taking goat milk (n=4 persons)				
<i>L. fermentum</i>	0	0	1	0 - 32,9

Statistically significant increase:

15 * The Fisher Exact Test showed the difference of counts in persons colonised with *Lactobacillus fermentum* ME-3 - $p < 0,0015$;

The Mann-Whitney Rank Sum Test showed the difference of relative share of *Lactobacillus fermentum* ME-3 in lactoflora.

20 Therefore, after consuming yoghurt in 3 weeks the microbe was present in the intestinal tract of all volunteers and the *Lactobacillus* sp. count was remarkably increased.

25

Table 6. The indices of oxidative stress of blood sera of volunteers (n=16) before and after consuming probiotic goat milk yoghurt during 3 weeks

Properties	Standard degree	Blood sera before trial	Blood trial after trial	Increase
TAA (LA-test, %)	36± 4.5	38 ± 3.5	45 ± 3.4	16%
TAS, mmol/L	1.2 ± 0.2	0.82 ± 0.14	1.14 ± 0.08	29%
Glutathione red-ox ratio (GSSG/GSH)	0.17±0.08	0.15 ± 0.01	0.11 ± 0.035	-32%
LDL lag-phase (time of resistance)	>30 min	41 ± 7.9	46 ± 8.6	11%
Basic value of diene conjugates (value of extinction)	< 0.3	0.27 ± 0.06	0.23 ± 0.06	-15%
Ox LDL (U/L)	>127	98±12	81 ±19	- 18%

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Explanations: LA-test - linolenic acid test; TAA - total anti-oxidative activity; TAS - total anti-oxidative status; GSSG - oxidized glutathione; GSH - reduced glutathione; GSSG/GSH - glutathione red-ox ratio, ox LDL - oxidized low-density lipoproteins.

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Thus all parameters determined in the blood sera of healthy volunteers changed beneficially during the 3-week yoghurt trial.

The application of the invention is not limited to the above-described example of achievement. In the range of the patent claim, some other variants of use are possible, for example the production of probiotic cheese and other milk products.

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CLAIM

The strain of micro-organism *Lactobacillus fermentum* ME-3 DSM
5 14241 as a novel anti-microbial and anti-oxidative probiotic
for use in pharmacy and food industry and in medicine as a
preparation resistant to antibiotics for the prophylaxis and
treatment of intestinal and uroinfections, also against
oxidative stress.

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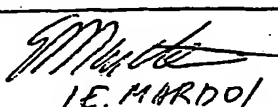
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Applicant's or agent's file reference	International application No. PCT/EE02/00006 PCT/EE02/00006
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Address of depositary institution (including postal code and country) Mascheroder Weg 1b D-38124 Braunschweig Germany	
Date of deposit 19.04.2001	Accession Number DSM 14241
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A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K35/74 C12N1/20 A61P31/04 A61P1/00

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Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K C12N A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

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C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	MIKELSAAR MARIKA ET AL: "Antagonistic and antioxidative activity of lactobacilli and survival in oxidative milieu." AMERICAN JOURNAL OF CLINICAL NUTRITION, vol. 73, no. 2S, February 2001 (2001-02), page 495S XP001105810 International Symposium on Probiotics and Prebiotics; Kiel, Germany; June 11-12, 1998 ISSN: 0002-9165 abstract	1
A	SEPP E ET AL: "Intestinal microflora of Estonian and Swedish infants." ACTA PAEDIATRICA, vol. 86, no. 9, 1997, pages 956-961, XP001105447 ISSN: 0803-5253 the whole document	1
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Date of the actual completion of the international search

23 September 2002

Date of mailing of the international search report

11/10/2002

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INTERNATIONAL SEARCH REPORT

International Application No
PCT/EE 02/00006

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	KULLISAAR TIIU ET AL: "Two antioxidative lactobacilli strains as promising probiotics." INTERNATIONAL JOURNAL OF FOOD MICROBIOLOGY, vol. 72, no. 3, 2002, pages 215-224, XP002214344 ISSN: 0168-1605 abstract table 1 page 223, column 1, last paragraph	1
P,A	MIKELSAAR MARIKA ET AL: "Intestinal lactobacilli of Estonian and Swedish children." MICROBIAL ECOLOGY IN HEALTH AND DISEASE, vol. 14, no. 2, June 2002 (2002-06), pages 75-80, XP001105478 June, 2002 ISSN: 0891-060X abstract table 2	1
P,A	ANNUK HEIDI ET AL: "Characterisation and differentiation of lactobacilli by lectin typing." JOURNAL OF MEDICAL MICROBIOLOGY, vol. 50, no. 12, December 2001 (2001-12), pages 1069-1074, XP002214345 ISSN: 0022-2615 abstract tables 2,3	1

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
9 January 2003 (09.01.2003)

PCT

(10) International Publication Number
WO 03/002131 A1

(51) International Patent Classification⁷: A61K 35/74,
C12N 1/20, A61P 31/04, 1/00

(21) International Application Number: PCT/EE02/00006

(22) International Filing Date: 21 June 2002 (21.06.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
P200100356 29 June 2001 (29.06.2001) EE

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(81) Designated States (national): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,
GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent
(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,
NE, SN, TD, TG).

Declarations under Rule 4.17:

- as to applicant's entitlement to apply for and be granted
a patent (Rule 4.17(ii)) for the following designations AE,
AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA,
CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES,
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KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG,
MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU,
SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG,
UZ, VN, YU, ZA, ZM, ZW, ARIPO patent (GH, GM, KE, LS,
MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent
(AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent
(AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU,
MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI,
CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)
- of inventorship (Rule 4.17(iv)) for US only

Published:

- with international search report
- with amended claims and statement
- with (an) indication(s) in relation to deposited biological
material furnished under Rule 13bis separately from the
description

Date of publication of the amended claims and statement:

6 March 2003

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: STRAIN OF MICRO-ORGANISM LACTOBACILLUS FERMENTUM ME-3 AS NOVEL ANTI-MICROBIAL AND ANTIOXIDATIVE PROBIOTIC

(57) Abstract: The strain of micro-organism *Lactobacillus fermentum* ME-3 is a novel anti-microbial and anti-oxidative probiotic. It has a high anti-microbial effect on *Escherichia coli*, *Shigella sonnei*, *Staphylococcus aureus*, *Salmonella typhimurium*, and moderate activity against *Helicobacter pylori* strains. The strain of micro-organism possesses Mn-superoxide dismutase and both its lysates and intact cells have high anti-oxidative activity, increasing the glutathione red-ox ratio in blood sera and able to capture toxic hydroxyl radicals. The strain of micro-organism could be used as a probiotic for the production of functional food (yoghurt, cheese) and non-comestibles (tablets, capsules) for the prophylaxis of intestinal and uroinfections, both for the prevention and treatment of chronic diseases, caused by prolonged oxidative stress.

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AMENDED CLAIMS

[received by the International Bureau on 06 December 2002 (06.12.02);
original claim 1 amended (1 page)]

The strain of micro-organism *Lactobacillus fermentum* ME-3 DSM 14241 as a novel anti-microbial and anti-oxidative probiotic for use in pharmacy and food industry and in medicine as a preparation resistant to antibiotics for the prophylaxis and treatment of gastrointestinal and uroinfections, also against chronic diseases induced by prolonged high-grade oxidative stress.

STATEMENT

The aim of present invention is to offer a strain of microorganism as a novel anti-microbial and anti-oxidative probiotic for use in pharmaceutical and food industry, also in medicine as an antibiotic resistant preparation for prophylaxis and treatment of gastrointestinal and uroinfections, also against oxidative stress (P.3, line 33-35, P.4, line 1-3)

Concerning the antimicrobial activity (bacteriostatic influence) of the object of invention, the strain *Lactobacillus fermentum* ME-3 expresses anti-microbial effect beside others on *Shigella sonnei*, *Salmonella typhimurium* ja *Helicobacter pylori* strains (Page 6, line 21-25; Page 7, Table 2). In addition, the property of *Lactobacillus fermentum* ME-3 to kill the food borne pathogens in milk (bacteriocidic effect) is firstly described (Page 7, Table 2 and line 6-11).

The innate resistance of *Lactobacillus fermentum* ME-3 against antimicrobial preparations (TMP-SMX, ofloxacin, aztreonam, cefoxitin and metronidazole) allows to use it as a preparation accompanying antibiotic treatment in case of gastrointestinal and uroinfections (Page 7, line 17-19; Page 8, line 1-3). This property has not been described elsewhere before.

The unique carbohydrate profile of the cell wall of *Lactobacillus fermentum* ME-3 enables to prevent the adhesion of uropathogenic *Escherichia coli* to the epithelial cells of

the upper urinary tract, a property that makes our strain applicable in the prophylaxis of urinary tract infections (Page 8, line 5-26) and has never been described before.

Concerning the antioxidative activity of the strain *Lactobacillus fermentum* ME-3 as the object of the present invention the different specific, principal and novel parameters were firstly described like expression of MnSOD, high-grade total antioxidative status (TAS, verified by internationally accepted method), principal parameters of glutathione (a signal molecule and central cellular antioxidant) system and the value of glutathione redox ratio (Page, 9 Table 3; Page 10, line 1-11, 15-19, 23-26).

Any antioxidativity (including antiatherogenicity) parameters found in human trials (*in vivo* trials) were not made public elsewhere. Therefore, only in this invention, an influence of consumption of ME-3 on human blood sera specific indices was described (Page 13-14, Table 6) and disclosed the appropriate numerical values. Actually, considering mainly these parameters (significant increase of TAS and oxygen resistance of LDL, lowering the level of oxidized LDL and its diene conjugates altogether indicate improvement of systemic antioxidativity and also significant lowering of cellular oxidative stress) it can be claimed that strain *Lactobacillus fermentum* ME-3 is a novel antioxidative (anti-atherogenic) probiotic (Page 11, line 25-30).

The persistence of the novel strain in gastrointestinal tract after consumption and the beneficial influence on the composition of the intestinal lactobacilli are described for the first time (Page 12, Table 5, line 14-22).

Thus the strain of microorganism *Lactobacillus fermentum* ME-3 (DSM 14241) figures a novel antimicrobial and antioxidative (anti-atherogenic) probiotic for use in pharmacy and food industry, and in medicine as a preparation resistant to some antimicrobials useful for the prophylaxis and as a preparation accompanying antibiotic therapy of gastrointestinal and urinary tract infections, also against chronic diseases (incl. atherosclerosis) induced by prolonged high-grade oxidative stress.